



Familial hypercholesterolemia

Familial hypercholesterolemia is an inherited condition characterized by very high levels of cholesterol in the blood. Cholesterol is a waxy, fat-like substance that is produced in the body and obtained from foods that come from animals (particularly egg yolks, meat, poultry, fish, and dairy products). The body needs this substance to build cell membranes, make certain hormones, and produce compounds that aid in fat digestion. In people with familial hypercholesterolemia, the body is unable to get rid of extra cholesterol, and it builds up in the blood. Too much cholesterol increases a person's risk of developing heart disease.

People with familial hypercholesterolemia have a high risk of developing a form of heart disease called coronary artery disease at a young age. This condition occurs when excess cholesterol in the bloodstream is deposited on the inner walls of blood vessels, particularly the arteries that supply blood to the heart (coronary arteries). The abnormal buildup of cholesterol forms clumps (plaques) that narrow and harden artery walls. As the plaques get bigger, they can clog the arteries and restrict the flow of blood to the heart. The buildup of plaques in coronary arteries causes a form of chest pain called angina and greatly increases a person's risk of having a heart attack.

Familial hypercholesterolemia can also cause health problems related to the buildup of excess cholesterol in tissues other than the heart and blood vessels. If cholesterol accumulates in the tissues that attach muscles to bones (tendons), it causes characteristic growths called tendon xanthomas. These growths most often affect the Achilles tendons, which attach the calf muscles to the heels, and tendons in the hands and fingers. Yellowish cholesterol deposits can develop under the skin of the eyelids and are known as xanthelasmata. Cholesterol can also accumulate at the edges of the clear, front surface of the eye (the cornea), leading to a gray-colored ring called an arcus cornealis.

Frequency

Familial hypercholesterolemia affects an estimated 1 in 200 to 1 in 250 people in most countries and is thought to be the most common inherited condition affecting the heart and blood vessels (cardiovascular disease). The condition occurs even more frequently in certain populations, including Afrikaners in South Africa, Lebanese, and Tunisians.

Causes

Mutations in the *APOB*, *LDLR*, *LDLRAP1*, or *PCSK9* gene cause familial hypercholesterolemia. Changes in the *LDLR* gene are the most common cause of this condition. The *LDLR* gene provides instructions for making a protein called a low-density lipoprotein receptor. This type of receptor binds to particles called low-density

lipoproteins (LDLs), which are the primary carriers of cholesterol in the blood. By removing LDLs from the bloodstream, these receptors play a critical role in regulating cholesterol levels. Some *LDLR* gene mutations reduce the number of low-density lipoprotein receptors produced within cells. Other mutations disrupt the receptors' ability to remove low-density lipoproteins from the bloodstream. As a result, people with mutations in the *LDLR* gene have very high levels of blood cholesterol. As the excess cholesterol circulates through the bloodstream, it is deposited abnormally in tissues such as the skin, tendons, and coronary arteries.

Less commonly, familial hypercholesterolemia is caused by mutations in the *APOB*, *LDLRAP1*, or *PCSK9* gene. Proteins produced from these genes are essential for the normal function of low-density lipoprotein receptors. Mutations in any of these genes prevent cells from making functional receptors or alter the receptors' function. Hypercholesterolemia results when low-density lipoprotein receptors are unable to remove cholesterol from the blood effectively. Some people with familial hypercholesterolemia do not have a mutation in one of these genes. In these cases, the cause of the condition is unknown.

Both genetic and environmental risk factors play roles in familial hypercholesterolemia. Lifestyle choices including diet, exercise, and tobacco smoking strongly influence the amount of cholesterol in the blood and the risk of coronary artery disease. Additional factors that impact the outcome of the condition include a person's gender, age, and health problems such as diabetes and obesity.

Familial hypercholesterolemia accounts for only a small percentage of all cases of high cholesterol. Researchers are working to identify and characterize additional genes that may influence cholesterol levels and the risk of heart disease in people with other forms of hypercholesterolemia.

Inheritance Pattern

Familial hypercholesterolemia resulting from mutations in the *LDLR*, *APOB*, or *PCSK9* gene have an autosomal dominant pattern of inheritance. Autosomal dominant inheritance means one copy of an altered gene in each cell is sufficient to cause the disorder. An affected person typically inherits one altered copy of the gene from an affected parent and one normal copy of the gene from the other parent.

Rarely, a person with familial hypercholesterolemia has a mutation in both copies of the *LDLR*, *APOB*, or *PCSK9* gene. This situation occurs when both parents have familial hypercholesterolemia, and each passes on one altered copy of the gene. The presence of two mutations results in a more severe form of familial hypercholesterolemia that usually appears in childhood.

When familial hypercholesterolemia is caused by mutations in the *LDLRAP1* gene, the condition is inherited in an autosomal recessive pattern. Autosomal recessive inheritance means the condition results from two altered copies of the gene in each cell. The parents of an individual with autosomal recessive hypercholesterolemia each carry

one copy of the altered gene, but their blood cholesterol levels are usually in the normal range.

Other Names for This Condition

- familial hypercholesterolaemia
- FH

Diagnosis & Management

Genetic Testing Information

- What is genetic testing?
[/primer/testing/genetictesting](#)
- Genetic Testing Registry: Familial hypercholesterolemia
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C0020445/>
- Genetic Testing Registry: Familial hypercholesterolemia 2
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1704417/>
- Genetic Testing Registry: Familial hypercholesterolemia 3
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1863551/>
- Genetic Testing Registry: Familial hypercholesterolemia 4
<https://www.ncbi.nlm.nih.gov/gtr/conditions/C1863512/>

Research Studies from ClinicalTrials.gov

- ClinicalTrials.gov
<https://clinicaltrials.gov/ct2/results?cond=%22familial+hypercholesterolemia%22>

Other Diagnosis and Management Resources

- GeneReview: Familial Hypercholesterolemia
<https://www.ncbi.nlm.nih.gov/books/NBK174884>
- Genomics Education Programme (UK)
<https://www.genomicseducation.hee.nhs.uk/documents/familial-hypercholesterolaemia/>
- MedlinePlus Encyclopedia: Familial hypercholesterolemia
<https://medlineplus.gov/ency/article/000392.htm>
- MedlinePlus Encyclopedia: High blood cholesterol and triglycerides
<https://medlineplus.gov/ency/article/000403.htm>
- MedlinePlus Encyclopedia: Xanthoma
<https://medlineplus.gov/ency/article/001447.htm>

- MedlinePlus Medical Tests: Cholesterol Levels
<https://medlineplus.gov/lab-tests/cholesterol-levels/>
- National Organization for Rare Disorders (NORD) Physician Guide
<https://rarediseases.org/physician-guide/homozygous-familial-hypercholesterolemia-hofh/>

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Familial hypercholesterolemia
<https://medlineplus.gov/ency/article/000392.htm>
- Encyclopedia: High blood cholesterol and triglycerides
<https://medlineplus.gov/ency/article/000403.htm>
- Encyclopedia: Xanthoma
<https://medlineplus.gov/ency/article/001447.htm>
- Health Topic: Cholesterol
<https://medlineplus.gov/cholesterol.html>
- Health Topic: High Cholesterol in Children and Teens
<https://medlineplus.gov/highcholesterolinchildrenandteens.html>
- Health Topic: Lipid Metabolism Disorders
<https://medlineplus.gov/lipidmetabolismdisorders.html>
- Medical Tests: Cholesterol Levels
<https://medlineplus.gov/lab-tests/cholesterol-levels/>

Genetic and Rare Diseases Information Center

- Familial hypercholesterolemia
<https://rarediseases.info.nih.gov/diseases/10416/familial-hypercholesterolemia>

Additional NIH Resources

- National Heart, Lung, and Blood Institute
<https://www.nhlbi.nih.gov/health-topics/blood-cholesterol>
- National Human Genome Research Institute
<https://www.genome.gov/Genetic-Disorders/Familial-Hypercholesterolemia>
- National Human Genome Research Institute: The Genomic Services Research Program (GSRP): Study of People with Unexpected Genetic Results
<https://www.genome.gov/Current-NHGRI-Clinical-Studies/Genomic-Services-Research-Program>

Educational Resources

- KidsHealth from the Nemours Foundation
<https://kidshealth.org/en/parents/cholesterol.html>
- MalaCards: hypercholesterolemia, autosomal dominant, 3
https://www.malacards.org/card/hypercholesterolemia_autosomal_dominant_3
- MalaCards: hypercholesterolemia, autosomal dominant, type b
https://www.malacards.org/card/hypercholesterolemia_autosomal_dominant_type_b
- MalaCards: hypercholesterolemia, autosomal recessive
https://www.malacards.org/card/hypercholesterolemia_autosomal_recessive
- Merck Manual Consumer Version
<https://www.merckmanuals.com/home/hormonal-and-metabolic-disorders/cholesterol-disorders/overview-of-cholesterol-and-lipid-disorders>
- Orphanet: NON RARE IN EUROPE: Heterozygous familial hypercholesterolemia
https://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=406

Patient Support and Advocacy Resources

- American Heart Association
<https://www.heart.org/en/health-topics/cholesterol>
- National Organization for Rare Disorders
<https://rarediseases.org/rare-diseases/familial-hypercholesterolemia/>

Clinical Information from GeneReviews

- Familial Hypercholesterolemia
<https://www.ncbi.nlm.nih.gov/books/NBK174884>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28Hypercholesterolemia,+Familial%5BMAJR%5D%29+AND+%28hypercholesterolemia%5BTI%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+360+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- HYPERCHOLESTEROLEMIA, FAMILIAL, 1
<http://omim.org/entry/143890>
- HYPERCHOLESTEROLEMIA, FAMILIAL, 2
<http://omim.org/entry/144010>

- HYPERCHOLESTEROLEMIA, FAMILIAL, 3
<http://omim.org/entry/603776>
- HYPERCHOLESTEROLEMIA, FAMILIAL, 4
<http://omim.org/entry/603813>

Sources for This Summary

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Citation on PubMed: <https://www.ncbi.nlm.nih.gov/pubmed/15199432>
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Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5457620/>
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Free article on PubMed Central: <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1421462/>
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